Platelet-Mediated Thrombosis and Drug-Eluting Stents

Juan F. Granada, MD; Matthew J. Price, MD; Patricia A. French, BS; Steven R. Steinhubl, MD; Donald E. Cutlip, MD; Richard C. Becker, MD; Susan S. Smyth, MD, PhD; Harold L. Dauerman, MD

Stent thrombosis is an example of device-induced, platelet-mediated arterial thrombosis. Rates of stent thrombosis can vary from <1% to >10% depending on the patient population, genetic predisposition, device type, pharmacological choices, and duration of antiplatelet pharmacotherapy. The Platelet Colloquium is an annual academic–industry–governmental think-tank meeting devoted to identifying research challenges in platelet biology and clinical applications. The latest meeting was held in Washington, DC, on January 25 to 26, 2011, and this review summarizes the discussions of biocompatible stent design, platelet function assessment, and prevention of thrombosis via short- and long-term P2Y12 platelet receptor antagonism.

Stent Design and Surface-Mediated Platelet Activation

The vascular injury induced by percutaneous coronary intervention (PCI) produces dynamic changes on the surface of human platelets. Activated platelets are among the first cells to arrive at the site of injury. Stent thrombosis results from the interaction of several procedural, anatomic, and genetically determined factors.

Early cellular and inflammatory events are influenced by the properties of the stent or its coating. First-generation drug-eluting stents (DES) used relatively thick struts and durable polymers. Research efforts focused on development of nonerodable biocompatible materials that could control the release of antiproliferative medications over several weeks. In vitro models showed that these devices appeared to be associated with increased platelet activation and adhesion compared with identical bare metal stents (BMS). The continuous presence of a durable polymer and drug has been postulated to be partly responsible for delayed arterial healing and enhanced stent thrombogenicity.

Second-generation DES modified some of these components by reducing strut thickness and polymeric drug load. In vitro data suggest that the lower polymeric drug load used in current everolimus-eluting stents may have a more favorable thrombogenic profile than BMS controls. Recent data also suggest that these devices might favorably affect inflammation and vascular healing after DES implantation. In clinical trials, second-generation DES appear to diminish some undesirable biological effects (thrombosis) seen with first-generation DES. This finding is supported by recent clinical trial data in the setting of ST-segment elevation myocardial infarction, suggesting that everolimus-eluting stents reduce the risk of late stent thrombosis with identical BMS controls in this high-risk population (EXAMINATION trial).

Further research in coating technologies has focused on bioerodable polymeric or polymer-free drug-releasing matrices, potentially allowing the drug-eluting platform to return to its bare metal backbone over several months. Several clinical studies have studied the safety and efficacy of third-generation DES using bioabsorbable coatings (Table 1) and polymer-free platforms (Table 2). These studies have reported very low rates of late stent thrombosis (LST), while maintaining long-term efficacy.

However, no randomized trial has shown a clear reduction in stent thrombosis with bioabsorbable versus durable polymers. This does not necessarily disprove the concept of durable polymer-induced adverse events; the rates of stent thrombosis may be too low to compare within a randomized trial. Thus, although the concept of complete polymer dissolution is attractive, questions about drug bioavailability, degradation profiles, and rebound inflammation remain.

One alternative strategy is to develop a drug elution vehicle that promotes healing and endothelialization. Although anti-CD34–coated stents have been shown to enhance stent coverage in vitro compared with sirolimus-eluting stents, several clinical studies using this technology have shown restenosis and LST rates comparable to those with other BMS platforms. Thus, a further step would be to promote endothe-
lialization by fixing antihuman-CD34 antibody to the DES surface. In a porcine model of coronary restenosis, anti-CD34 antibody-coated sirolimus-eluting stents were associated with greater endothelialization at 3 and 14 days, compared with conventional sirolimus-eluting stents. However, a clear clinical benefit of stents coated with this technology has not been shown. A randomized clinical trial using this dual approach (prohealing and sirolimus elution) is under development.

Preclinical modeling continues to provide meaningful insights regarding the potential for next-generation DES to improve clinical outcomes. For example, bench testing of stent thrombogenicity, combined with computational modeling, appears to correlate with clinical outcomes seen in large randomized trials of second- versus first-generation DES. If second-generation DES with durable polymers continue to produce excellent safety profiles, showing the additional value of newer technologies (ie, polymer-free coatings) or fourth-generation DES (bioabsorbable polymers on bioresorbable scaffolds) will become very difficult.

The concept of a fourth-generation, fully bioresorbable polylactide-everolimus DES is especially attractive as a potential way to restore vasomotion and endothelial function to potentially limit any hazard of LST. The challenge will be to show superiority to second- and third-generation DES with respect to major adverse cardiovascular events (MACE) or softer end points (measures of healing or endothelial function). Based on likely low rates of clinical events in these future trials,

<table>
<thead>
<tr>
<th>Stent Type (Manufacturer)</th>
<th>Drug</th>
<th>Stent Material</th>
<th>Polymer Type</th>
<th>Study Type (No. of Patients)</th>
<th>In-Stent Late Loss, mm</th>
<th>Binary Restenosis, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>CoStar (Conor Medical)†²³</td>
<td>Paclitaxel</td>
<td>CoCr</td>
<td>PLGA</td>
<td>Randomized controlled trial (CoStar n=989 vs Taxus n=686)</td>
<td>0.64 vs 0.26*</td>
<td>17.9 vs 4.1*</td>
</tr>
<tr>
<td>Supralimus (Sahajanand Medical)†⁴</td>
<td>Sirolimus</td>
<td>SS</td>
<td>PLLA PLGA, PLC, PVP</td>
<td>First in man n=100</td>
<td>0.09</td>
<td>0.0</td>
</tr>
<tr>
<td>Excel Stent (AW Medical System)†⁵</td>
<td>Sirolimus</td>
<td>SS</td>
<td>PLA</td>
<td>Registry n=2077</td>
<td>0.21</td>
<td>3.8</td>
</tr>
<tr>
<td>NEVO (Cordis)†⁶</td>
<td>Sirolimus</td>
<td>CoCr</td>
<td>PLGA Reservoirs</td>
<td>Randomized controlled trial Nevo (n=202 vs PES n=192)</td>
<td>0.13 vs 0.36*</td>
<td>1.1 vs 8.0*</td>
</tr>
<tr>
<td>BioMatrix (Biosensors)†²³</td>
<td>Biolimus A9</td>
<td>SS</td>
<td>Abluminal PLA</td>
<td>Randomized controlled trial BES (n=857 vs SES n=850)</td>
<td>0.13 vs 0.19</td>
<td>20.9 vs 23.3*</td>
</tr>
<tr>
<td>NOBORI (Terumo)†⁸</td>
<td>Biolimus A9</td>
<td>SS</td>
<td>Abluminal PLA</td>
<td>Randomized controlled trial BES (n=153 vs SES n=90)</td>
<td>0.11 vs 0.32*</td>
<td>0.7 vs 6.2†</td>
</tr>
<tr>
<td>SYNERGY (Boston Scientific; JACTAX Liberté (Boston Scientific)†²¹</td>
<td>Everolimus</td>
<td>PtCr</td>
<td>PLGA Rollcoat Abluminal</td>
<td>Randomized controlled trial SD vs (LD vs PROMUS Element n=291)</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Combi EPC+ drug (OrbusNeich; NCT00967902)</td>
<td>Sirolimus</td>
<td>SS</td>
<td>Abluminal PLA</td>
<td>Randomized controlled trial Combi (stent vs PES; n=180)</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Elixir Myolimus (Elixir Medical)†¹⁹</td>
<td>Myolimus</td>
<td>CoCr</td>
<td>Abluminal PLA</td>
<td>First in man n=15</td>
<td>0.15</td>
<td>0</td>
</tr>
<tr>
<td>Infinnium (Sahajanand)†²₀</td>
<td>Paclitaxel</td>
<td>SS</td>
<td>PLLA PLGA, PLC PVP</td>
<td>Randomized controlled trial Infinnium (n=111 vs BMS n=57)</td>
<td>0.54 vs 0.90†</td>
<td>8.3 vs 25.5*</td>
</tr>
<tr>
<td>JACTAX Liberté (Boston Scientific)†²¹</td>
<td>Paclitaxel</td>
<td>SS</td>
<td>JAC polymer</td>
<td>First in man n=103</td>
<td>0.33</td>
<td>5.2</td>
</tr>
</tbody>
</table>

BES indicates biolimus-eluting stent; BMS, bare metal stent; CoCr, cobalt chromium; EPC, endothelial progenitor cell; JAC, juxtaposed abulminal coating; LD, low dose; NA, not available; PES, paclitaxel-eluting stent; SES, sirolimus-eluting stent; SS, stainless steel.

Table 2. Balloon-Expandable Stents Using Polymer-Free DES Platforms

<table>
<thead>
<tr>
<th>Stent Type (Manufacturer)</th>
<th>Drug</th>
<th>Stent Material</th>
<th>Delivery Method</th>
<th>Study Type No. of Patients</th>
<th>In-Stent Late Loss, mm</th>
<th>Binary Restenosis, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>AmazoniaPax (Minvasys)²²</td>
<td>Paclitaxel</td>
<td>CoCr</td>
<td>Abluminal microspray crystallization process</td>
<td>First in man Pax n=16 vs PES n=15</td>
<td>0.77 vs 0.42</td>
<td>NA</td>
</tr>
<tr>
<td>BioFREEDOM (Biosensors)²²</td>
<td>Biolimus A9 (SD and LD)</td>
<td>SS</td>
<td>Microporous surface</td>
<td>First in man SD n=25 vs LD n=25 vs PES n=25</td>
<td>0.08 vs 0.37*</td>
<td>NA</td>
</tr>
<tr>
<td>VESTA sync (MIV Therapeutics)²⁴</td>
<td>Sirolimus</td>
<td>SS</td>
<td>Nanoporous hydroxyapatite</td>
<td>First in man n=15</td>
<td>0.36</td>
<td>0</td>
</tr>
<tr>
<td>Yukon (Translumina)²⁵</td>
<td>Rapamycin</td>
<td>SS</td>
<td>Microporous surface</td>
<td>Randomized controlled trial Yukon n=225 vs PES n=225</td>
<td>0.48 vs 0.48</td>
<td>12.6 vs 11.6</td>
</tr>
</tbody>
</table>

CoCr indicates cobalt chromium; DES, drug-eluting stents; LD, low dose; NA, not available; PES, paclitaxel-eluting stent; SD, standard dose; SS, stainless steel.

*P<0.001.
†P<0.05.
pooling of several randomized studies likely will be needed to evaluate the long-term efficacy and safety of emerging technologies, including the practical question of whether new designs will allow shorter courses of dual antiplatelet therapy (DAPT).

In Vivo Testing With Platelet Function Assays

Currently, all DES aim to prevent surface-mediated platelet activation through at least 12 months of DAPT—aspirin and a P2Y12 receptor antagonist. Clopidogrel is the most commonly used P2Y12 antagonist, but its pharmacodynamic effects are variable. As a result, various platelet function tests have been proposed to monitor and guide DAPT in this setting. One potential mechanism for limiting the risk of platelet activation and device thrombosis is to individualize the pharmacological approach according to patient-mediated (not device-mediated) risk.

Previous studies in this regard are limited in several important ways. First, the cutoff values derived from the studied populations (primarily at single centers) were not prospectively confirmed in independent validation cohorts. Further, multivariable models showing independent associations between on-treatment reactivity (OTR) while receiving clopidogrel and outcomes were likely “overfitted”; they included too many covariates for too few events. Finally, these studies did not address whether OTR truly is a modifiable risk factor for future cardiovascular events.

Table 3 shows completed and ongoing randomized studies of individualized antiplatelet therapy during or after percutaneous coronary intervention (PCI).29,30 Gauging Responsiveness with A VerifyNow assay-Impact on Thrombosis And Safety (GRAVITAS; NCT00645918) was designed to assess whether high-dose clopidogrel would be superior to standard-dose clopidogrel in preventing MACE at 6 months among patients with high OTR after DES implantation.29 Of 5429 patients screened with the VerifyNow P2Y12 test after PCI, 2214 (41%) had high OTR while receiving clopidogrel, defined as >230 P2Y12 reaction units. They were randomized to receive either a 75-mg daily maintenance dose of clopidogrel, or another 600-mg loading dose, followed by 150-mg daily maintenance dosing.

The incidence of MACE at 6 months (primary end point) was 2.3% in both groups (hazard ratio [HR], 1.01; 95% CI, 0.58 to 1.76; \( P=0.97 \)).29 Stent thrombosis developed in 0.5% of patients in the higher-dose group and 0.7% of the standard-dose group (\( P=0.42 \)). Bleeding rates did not differ significantly, although the proportions of patients with persistently high OTR were modestly but significantly reduced from baseline at 30 days and 6 months.

A separate observational analysis compared GRAVITAS patients assigned to standard-dose clopidogrel after PCI by the presence (n=1105) or absence (n=586) of persistently high OTR.31 Patients with high OTR had a nonsignificantly higher rate of MACE than patients without high OTR (HR, 1.68; 95% CI, 0.76 to 3.72; \( P=0.20 \)).29 In post hoc analysis, patients with lower levels of OTR after PCI or during follow-up had a significantly lower risk of MACE.31

A more recent trial illustrates the difficulty of showing differences between therapies when event rates are low. In July 2009, the Testing platelet Reactivity In patients undergoing elective stent placement on clopidogrel to Guide alternative thERapy with prasugrel (TRIGGER-PCI; NCT00910299) began enrollment of its 2150 expected patients with stable coronary artery bypass surgery undergoing successful, elective PCI with DES. On March 18, 2011, the sponsor stopped the study after a preliminary, blinded analysis of the first 250 patients to complete follow-up revealed that the trial would not generate enough primary end point events (cardiovascular death or myocardial infarction [MI] at 6 months) for analysis. The study had been designed assuming a 7% incidence of the primary end point for this interval.

Although GRAVITAS did not support treatment with high-dose clopidogrel when 1 platelet function test identified...
high OTR after PCI, it did illustrate some important phenomena. First, OTR was shown to be dynamic for the first month after PCI. Second, the pharmacodynamic effect of the higher maintenance dose was marginal relative to standard dosing in patients with high OTR. Third, the 6-month MACE rate was relatively low with modern DES and techniques used in patients with stable coronary artery bypass surgery. Therefore, very large cohorts will be required to show independent associations between OTR and outcomes, given the multitude of clinical predictors of high OTR and the infrequency of events.

Demonstrating a benefit of antiplatelet therapy tailored to platelet function will be similarly challenging, given the large sample sizes required to provide adequate power to detect outcome differences between treatment groups. Ongoing randomized clinical trials, Thrombocyte Activity Reassessment and GEneTyping for PCI (TARGET-PCI; n=1500; NCT01177592) and Assessment with a double Randomization of (1) a fixed dose versus a monitoring-guided dose of aspirin and Clopidogrel after DES implantation, and (2) Treatment Interruption versus Continuation, 1 year after stenting (ARCTIC; n=2500), are likely underpowered in this regard. In addition, given the low event rates observed in stable coronary artery bypass surgery patients, the net clinical benefit of potent P2Y12 inhibitors in patients with high OTR likely will be narrow. Future studies should focus on larger, “enriched” patient populations (eg, acute coronary syndromes) and longer follow-up. For now, the roles of platelet function testing to determine patient-mediated risk and to tailor antiplatelet therapy to prevent platelet-mediated device thrombosis remain unproven.

The American College of Cardiology/American Heart Association guidelines state that physicians may consider platelet function testing to determine platelet inhibitory response in patients receiving thienopyridine therapy if the results of testing might alter management (Class IIb, level of evidence, B). Similarly, the European Society of Cardiology guidelines state that platelet function testing may be considered in selected cases when clopidogrel is used (Class IIb, level of evidence, B).

Pharmacological Choices and Platelet Activation in DES Thrombosis

In the early stent experience, 5-drug antithrombotic regimens were not uncommon and included aspirin, dipyridamole, dextran, and prolonged heparin followed by warfarin. Despite this intensive therapy, stent thrombosis rates were routinely at or above 5%, with major bleeding rates 2 to 4 times higher. Several randomized trials showed substantial decreases in thrombotic complications with aspirin plus thienopyridine therapy versus either aspirin alone or aspirin plus warfarin (Figure 1). These studies highlighted the critical role of using inhibitors of platelet activation (aspirin and P2Y12 receptor antagonists) to prevent stent thrombosis. The fact that DAPT not only was a more effective antithrombotic but also significantly reduced bleeding risk compared with a warfarin-based regimen is an important lesson; there needs to be a tradeoff of increased bleeding for improved thrombosis prevention when the correct thrombotic pathway is targeted.

Platelet activation occurs almost immediately after stenting and appears to peak ≈2 to 4 hours afterward. In addition to aspirin, platelet P2Y12 antagonists have been the primary agents used to minimize platelet activation. Although clopidogrel has been the gold standard, it has several limitations. Specifically, it was never designed to maximize inhibition of the platelet P2Y12 receptor; in fact, it is thought to routinely inhibit only ≈80% of the P2Y12 receptors on the platelet surface. In contrast, the newer thienopyridine prasugrel and the nonthienopyridine ticagrelor were specifically designed to achieve higher levels of inhibition. These agents appear to block 100% of platelet surface P2Y12 receptors. This enhanced antiplatelet effect is likely explained by the more efficient generation of prasugrel’s active metabolite compared with clopidogrel’s.

Similarly, ticagrelor is a nonthienopyridine P2Y12 receptor antagonist with greater potency and more rapid achievement of therapeutic levels compared with clopidogrel. The platelet inhibition and patient outcomes trial showed a significant 16% reduction in the primary end point (death from vascular causes, MI, or stroke) and reductions in stent thrombosis and all-cause mortality with ticagrelor compared with clopidogrel.
in patients with acute coronary syndromes. Prasugrel and ticagrelor have different pharmacokinetics and side effect profiles, but both agents increase non-coronary artery bypass surgery related bleeding compared with clopidogrel.42 The benefit of both prasugrel and ticagrelor with respect to prevention of stent thrombosis has led to US (prasugrel)32 and European (ticagrelor and prasugrel)33 Class I guideline recommendations for these more potent agents. Thus, the clinical trial data are consistent with the concept that higher levels of P2Y₁₂ inhibition translate into greater protection from thrombotic events compared with clopidogrel.52,53

Clinical trials of newer P2Y₁₂ inhibitors and novel inhibitors of other platelet thrombin receptors (such as protease activated receptor-1) offer the possibility of enhanced inhibition of platelet activation in various clinical scenarios. Cangrelor and elinogrel are both intravenous nonthienopyridine P2Y₁₂ inhibitors (elinogrel also has an oral form). In vitro and ex vivo data show that they provide high levels of inhibition against adenosine diphosphate-induced platelet aggregation, but Phase 3 trials of cangrelor were disappointing.44 Elinogrel is now entering Phase 3 study, after earlier clinical trials showed promising results.45 Protease activated receptor-1 inhibitors offer a novel mechanism for minimizing platelet activation with encouraging Phase 2 data; however, Phase 3 data have yet to be published.

### Blocking Platelet-Mediated Thrombosis Over the Long Term

Although devices, pharmacological therapies, patient testing strategies, and development have focused on the general prevention of device-mediated thrombosis, the specific issue of LST has been most readily addressed by extending the duration of PCI pharmacotherapies. Whether new biocompatible/nondurable polymer DES will obviate this need remains to be determined. For the immediate future, the question is not whether extended DAPT is needed to prevent late platelet-mediated thrombosis, but for how long.

Early studies of sirolimus-eluting stents and paclitaxel-eluting stents mandated 2 or 6 months of DAPT.50,51 In 2006, however, registry data raised concern about the risk of LST-related death and MI.52,53 A prospective cohort study also had identified premature DAPT discontinuation as an independent predictor of stent thrombosis within the first 9 months among 2229 patients who had been prescribed DAPT for 3 to 6 months after DES implantation.54 Another large cohort study reported that LST can occur at an annual rate of 0.6% up to 3 years after DES implantation.55 Long-term concern about LST has been highlighted by findings from the recent SIRolimus-eluting versus paciTXel-eluting stents for coronary revascularization (SIRTAx) LATE trial, in which the annual rate of stent thrombosis was 0.65% between 1 and 5 years after implantation of first-generation DES.56,57 Although careful examination of patient-level data has refuted initial fears of increased mortality after DES implantation, concerns remain about the risk of LST beyond 1 year.

Current guideline recommendations reflect ongoing uncertainty: one set of recommendations calls for 12 months of DAPT after placement of a DES,68 and the other recommends 6 to 12 months.69 Clinical practice also varies considerably. In 1 US-based trial comparing everolimus- and paclitaxel-based DES, compliance with DAPT at 2 years was 69%,60 whereas a European study of the same 2 stents revealed a 15% compliance rate at 2 years.61 Contrary to US practice, a European Society of Cardiology statement on bleeding complications after PCI recommends that use beyond 12 months after DES placement be the exception, not routine practice.62 Registry data have not shown a clear benefit for extending DAPT beyond 6 months (Table 4).63–70 in all patients treated with DES. Clarification of the benefits of 6 versus 12 months of DAPT in stable PCI patients is being investigated in the Intracoronary Stenting and Antithrombotic Regimen: Safety and Efficacy of a 6-month DAT After Drug-eluting Stenting trial (NCT00661206).

In 2006, the US Food and Drug Administration agreed that at least 12 months of DAPT should be recommended for “off-label uses of DES.”71 Simultaneously, the Food and Drug Administration began asking DES manufacturers for studies of the optimal duration of DAPT as a condition of approval. The Food and Drug Administration’s Critical Path Initiative created a public–private collaboration, the DAPT

### Table 4. Registries Assessing Outcomes Relative to DAPT Adherence

<table>
<thead>
<tr>
<th>Study</th>
<th>n</th>
<th>Population</th>
<th>Study Duration</th>
<th>Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Airoldi63</td>
<td>3021</td>
<td>DES</td>
<td>18 mo</td>
<td>DAPT protective for ST only during first 6 mo; median time to ST with DAPT cessation &lt;6 mo = 14 d; 90 d thereafter</td>
</tr>
<tr>
<td>Park64</td>
<td>2873</td>
<td>Event-free DES</td>
<td>3 y</td>
<td>No reduction in death, MI, ST with DAPT after 1 y</td>
</tr>
<tr>
<td>j-Cypher65</td>
<td>10 778</td>
<td>SES</td>
<td>2 y</td>
<td>Discontinuation of DAPT, but not of aspirin alone, associated with ST at any time points</td>
</tr>
<tr>
<td>Schultz66</td>
<td>6816</td>
<td>DES</td>
<td>4 y</td>
<td>DAPT protective for ST only during first 6 mo; median time to ST with DAPT discontinuation &lt;6 mo = 9 d; 104 d thereafter</td>
</tr>
<tr>
<td>Roy67</td>
<td>2889</td>
<td>DES</td>
<td>1 y</td>
<td>DAPT not protective for ST after 6 mo</td>
</tr>
<tr>
<td>Van Werkum68</td>
<td>21 009</td>
<td>DES, BMS</td>
<td>30.9 mo</td>
<td>ST associated with DAPT discontinuation within 1 y; risk greatest if discontinued &lt;30 d</td>
</tr>
<tr>
<td>Petersen69</td>
<td>9256</td>
<td>DES</td>
<td>1 y</td>
<td>Prolonged DAPT use associated with greater bleeding risk but lower risk of death or nonfatal MI</td>
</tr>
<tr>
<td>e-SELECT70</td>
<td>15 147</td>
<td>SES</td>
<td>1 y</td>
<td>ST associated with any DAPT discontinuation within 30 d</td>
</tr>
</tbody>
</table>

BMS indicates bare metal stent; DAPT, dual antiplatelet therapy; DES, drug-eluting stents; MI, myocardial infarction; ST, stent thrombosis.
study (NCT00977938), to address this issue.72 Four device manufacturers, as well as international government and academic centers, are conducting a randomized, placebo-controlled trial of DAPT (clopidogrel or prasugrel with aspirin) among 20,645 patients who are (1) free from death, MI, stroke, repeat coronary revascularization, major bleeding, or stent thrombosis 12 months after implantation of a DES or BMS, and (2) already compliant with 12 months of DAPT after stent implantation. Eligible patients will be randomized to receive another 18 months of DAPT or begin taking placebo with aspirin. The primary end points are the composite incidence of death, MI, or stroke up to 33 months after stent implantation, stent thrombosis over the same interval, and major bleeding over this interval.

Randomization to prolonged versus 12-month DAPT duration in the DAPT study is expected to be complete in May 2012, and results are expected to be available in the spring of 2014. The study already has had to overcome some challenges, however. For example, the trial design assumed that 80% of patients would be eligible for randomization, but only 60% have satisfied all criteria and agreed to randomization to date. Two further caveats: the study data cannot answer questions about outcome differences by stent type (first- versus second-generation DES) or individual thienopyridine agent, and the trial will have only limited ability to assess the relation between short-term interruptions in DAPT and outcomes.

In the meantime, current practice may be best defined by large registry data (Table 4).63–70 Most of these analyses identified a protective benefit of DAPT during the first 6 months after DES implantation, but the data conflict regarding subsequent benefit (Figure 2).63 In addition, benefits at all time points must always be weighed against the potential increase in bleeding risk (Figure 3).69 Finally, type and duration of DAPT therapies are not the only issues pertaining to this debate; bleeding risk and anti-thrombotic benefits of DAPT might vary according to aspirin dose69,74 or regional variations in care.75

Once we reach 2014, will we no longer need to rely on postapproval surveillance data and registry studies to determine the optimal duration of DAPT to prevent coronary stent thrombosis? This seems unlikely; pharmacological agents, polymers, stent designs, and practice patterns continue to evolve far more rapidly than does the randomized evidence. It is unlikely that we can perform 20,000-patient clinical trials for each new generation of DES, and it might be false to assume a class effect with respect to device-mediated thrombosis. Thus, preclinical testing and assessment of patient-related risk will continue to be critical. Well-organized registries can continue to offer useful information in this regard, keeping in mind the observational nature of the data collected.

Conclusions

In summary, stent thrombosis cannot be viewed from a singular device, patient, or pharmacological perspective; prevention of platelet-mediated stent thrombosis requires a 3-fold approach focusing on the development of improved polymer systems, assessment of individual patient-related platelet pathophysiology, and optimization of the dose and duration of platelet receptor antagonist therapy:

- Stent thrombosis is an example of acute and delayed device-induced, platelet-mediated device thrombosis.
- Emerging DES platforms are moving toward biocompatible and bioerodible polymers, which aim to prevent potential platelet activation and inflammation associated with early earlier-generation DES designs.
- Although comparative clinical trials will provide an important evidence base, distinguishing the potential advantages of iterative changes in DES design will likely hinge on in vitro testing methods, soft end points (stent healing, endothelial function), and well-designed registry analyses.
- The scientific community still faces the challenges of treating patient-specific risks of platelet-mediated stent thrombosis with point-of-care testing.
- DAPT focusing on P2Y12 receptor antagonism has a proven role in preventing platelet-mediated stent thrombosis. The role of newer agents in preventing stent thrombosis is proven, but the optimum treatment duration of DAPT therapy remains uncertain.

Sources of Funding

The 2011 Platelet Colloquium and development of this manuscript were supported by unrestricted grants from Abbott Vascular, Inc; Redwood...
Disclosures

Dr Granada has received grants from Abbott Vascular, Boston Scientific, and Medtronic. Dr Price has received grants from Accuminetics, BMS/sanofi-aventis, and Quest Diagnostics; has had speaker’s bureau appointments for Daiichi/Lilly; has received honoraria from BMS/sanofi-aventis, Medtronic, St. Jude Medical, Boston Scientific, and AstraZeneca; and has consulted for Accuminetics, BMS/sanofi-aventis, Daiichi/Lilly, and Medicare. Ms. French has consulted for Regado Biosciences. Dr Steinlibuh is a former employee of The Medicines Company. Dr Cutlip has received grants from Medtronic and has consulted for AstraZeneca. Dr Becker has received grants from AstraZeneca, Bayer, and Regado Biosciences; and has consulted for Daiichi/Lilly and Boehringer-Ingelheim. Dr Smyth has received grants from AstraZeneca and The Medicines Company, and has consulted for BMS/sanofi-aventis. Dr Dauerman has consulted for Abbott Vascular, Medtronic, and MDS Scientific; has consulted for Abbott Vascular, Medtronic, MDS Scientific, Novartis, The Medicines Company, Gilead, and St. Jude Medical; and has served as an expert witness for the defense in New Hampshire.

References


Key Words: platelets; stents; blood flow; thrombosis; antiplatelet agents.
Supplemental Material

Appendix. Participants in the 2011 Platelet Colloquium

Richard C. Becker, MD, Duke Clinical Research Institute, Durham, NC; Danny Bluestein, PhD, State University of New York at Stony Brook; Christopher P. Cannon, MD, Harvard Medical School, Boston, MA; Mack Consigny, PhD, MBA, Abbott Vascular, Inc., Santa Clara, CA; Donald E. Cutlip, MD, Harvard Medical School, Boston, MA; Harold L. Dauerman, MD, University of Vermont, Burlington; Michael J. Eppihimer, PhD, Boston Scientific Corporation, Natick, MA; Andrew Farb, MD, U.S. Food and Drug Administration, Silver Spring, MD; Alok Finn, MD, Emory University, Atlanta, GA; Jane E. Freedman, MD, Boston University School of Medicine, Boston, MA; Patricia A. French, Left Lane Communications, Chapel Hill, NC; Gaurav Girdhar, PhD, State University of New York at Stony Brook; Juan F. Granada, MD, Cardiovascular Research Foundation, Orangeburg, NY; Peter L. Gross, MD, MSc, McMaster University, Hamilton, Ontario, Canada; Willibald Hochholzer, MD, Harvard Medical School, Boston, MA; Mary V. Jacoski, MS, Boston Scientific Corporation, Marlborough, MA; Reema Jasuja, PhD, Beth Israel Deaconess Medical Center, Boston, MA; Lisa K. Jennings, PhD, University of Tennessee Health Science Center, Memphis; Aditee Kurane, PhD, St. Jude Medical, St. Paul, MN; Donald R. Lynch, Jr., MD, Johns Hopkins Hospital, Baltimore, MD; Robert Melder, ScD, Medtronic Cardiovascular, Santa Rosa, CA; Jayne Prats, PhD, The Medicines Company, Waltham, MA; Matthew J. Price, MD, Scripps Translational Science Institute, La Jolla, CA; Jesse W. Rowley, PhD, University of Utah, Salt Lake City; Maurice Rozek, MD, Daiichi Sankyo, Inc., Parsippany, NJ; Christopher P. Rusconi, PhD, Regado Biosciences, Inc., Durham, NC; Alec Sheehy, PhD, Abbott Vascular Inc., Santa Clara, CA; Susan S. Smyth, MD, PhD, University of Kentucky, Lexington; Steven R. Steinhubl, MD, Geisinger Health System, Danville, PA; Fanmuyi Yang, University of Kentucky, Lexington; Guy A. Zimmerman, MD, University of Utah, Salt Lake City.